All-optical production of ^{7}Li Bose-Einstein condensation using Feshbach resonances.

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We show an all-optical method of making ^{7}Li condensate using tunability of the scattering length in the proximity of a Feshbach resonance. We report the observation of two new Feshbach resonances on $|F = 1, m_F = 0\rangle$ state. The narrow (broad) resonance of 7 G (34 G) width is detected at 831 ± 4 G (884^{+4}_{-13} G). Position of the scattering length zero crossing between the resonances is found at 836 ± 4 G. The broad resonance is shown to be favorable for run away evaporation which we perform in a crossed-beam optical dipole trap. Starting directly form the phase space density of a magneto-optical trap we observe a Bose-Einstein condensation threshold in less than 3 s of forced evaporation.

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I. INTRODUCTION

Achieving quantum degeneracy in ultracold atomic gases by all-optical means becomes a well accepted experimental technique because of several inherent advantages [1, 2, 3, 4, 5, 6, 7]. Optical traps allow strong confinement resulting in high collision rates and rapid evaporative cooling. Confinement of arbitrary spin states and spin state mixtures are readily obtained. The possibility to tune interactions via Feshbach resonances usually requires optical trapping as they frequently occur in states that cannot be trapped magnetically [8, 9]. Finally, large current coils needed for magnetic-field trapping that restrict optical access are avoided.

The first successful demonstration of an all-optically achieved ^{87}Rb Bose-Einstein condensation (BEC) allowed significant increase in the rate of BEC production and the resulting condensates were F=1 spinors [1]. However, the main driving force behind the search for all-optical techniques was the need to condense specific atoms where the 'conventional' evaporation in the magnetic trap was not possible. Two prominent examples are spinless, and thus magnetically untrappable, BEC of Yb atoms achieved in a doubled YAG crossed dipole trap [2] and a BEC of ^{133}Cs atoms [3] for which the strongly enhanced two-body losses from the magnetically trappable states prevent the condensate formation in the 'standard' way [10].

Although ^{7}Li atoms can be evaporatively cooled in a magnetic trap [11], the task remains challenging due to several reasons. First, ^{7}Li atoms posses a relatively small scattering length ($a = -27a_0$, where a_0 is the Bohr radius) and high two-body loss rate [12]. Second, the initial phase space density is unfavorably limited by the absence of polarization-gradient cooling mechanism. Third, since the scattering length drops with increased temperature and crosses zero at $T = 8$ mK [13], the use of adiabatic compression to increase the elastic collisional rate is ineffective. Therefore, the strong magnetic confinement needed to keep evaporation time comparable with heavier alkalies, such as Rb and Na, requires the design of a miniaturized trap. This is done by either a small-volume vacuum chamber and high currents [14] or a vacuum compatible minitrap [15] which both increase the experimental complexity. Finally, even if the strong confinement is achieved the scattering length is still negative which prevents the formation of a stable BEC.

In this paper we show an all-optical way to condense ^{7}Li atoms using tunable interatomic interactions. We use a 100 W Ytterbium fiber laser to produce $\sim$2 mK potential well which traps $\sim 10^6$ atoms from a magneto-optical trap (MOT). We explore two new Feshabach resonances on $|F = 1, m_F = 0\rangle$ state and find that one of them is favorable for the efficient forced evaporation starting directly from the phase space density achieved in the MOT. We obtain a BEC on $|F = 1, m_F = 0\rangle$ internal state in less than 3 s of evaporation time.

II. EXPERIMENTAL DETAILS

The ^{7}Li atoms' route to quantum degeneracy starts in an oven where they are heated up to 450^0C to increase their vapor pressure. An atomic beam is collimated by two distant apertures and slowed down in an increased magnetic-field Zeeman-slower. The capture velocity of the slower is set to $\sim$800 m/s.

Our MOT design and parameters are similar to that described in ref [16]. After a loading time of 3 s, the trap contains $\sim 10^9$ atoms at a temperature of 1.2 mK. Trap life-time, limited by vacuum, is measured to be 10 s. The MOT parameters, such as pump and repump detunings and magnetic field gradient, are optimized to maximize the number of atoms. To improve the initial phase space density we implement a so called compressed MOT (CMOT) stage [16, 17]. For 50 ms the laser intensities are attenuated, detunings are decreased and the magnetic field gradient is increased. As a result the temperature is reduced to 300 μK and nearly half of the atoms are lost. By the end of this phase $\sim 5 \times 10^8$ atoms are left in the trap with density of $n = 4 \times 10^{11}$ atoms/cm^3 and phase-space density of $\rho = 2 \times 10^{-5}$.

The realization of the optical-dipole trap is shown in Fig. 1. A CW Ytterbium fiber laser generates 100 W of linearly polarized light at 1.07 μm. The first order

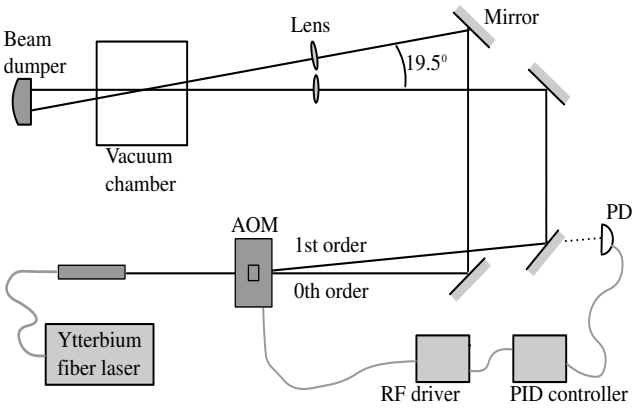

Figure 1: A schematic representation of the optical-dipole trap experimental setup. The first order diffraction beam is focused to a waist of 31 μm and aligned to the MOT center. The zeroth order beam is weakly focused to 300 μm. It crosses the first order beam at an angle of 19.5^0 and it only becomes important at the final stages of evaporation. AOM - Acousto-optic modulator; PD - Photodetector.

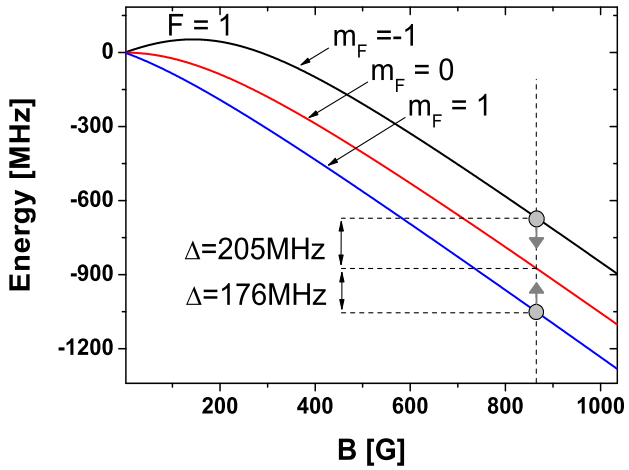

Figure 2: Zeeman splitting of the $|F = 1\rangle$ hyper-fine state of ^{7}Li atoms. With the presence of high magnetic fields, spin-flip collisions between $|m_F = -1\rangle$ and $|m_F = 1\rangle$ states result in their transfer into $|m_F = 0\rangle$ with an excess of kinetic energy equal to 29 MHz (~ 1.4 mK).

diffraction beam of an acousto-optic modulator (AOM), with 80% diffraction efficiency, is focused at the center of the MOT to a waist of 31 μm. The zero order diffraction beam is only weakly focused to 300 μm and it crosses the first order beam at an angle of 19.5^0. When maximum RF power is applied to the AOM this beam is very weak and it has no effect on the atoms. At full laser power (65 W in the trap region) the trap depth, produced by the first order diffraction beam, is estimated to be ~ 2 mK. Oscillation frequencies in the trap are predicted to be $\omega_r = 2\pi \times 16$ kHz and $\omega_z = 2\pi \times 120$ Hz for the radial and longitudinal directions, respectively. The radial oscillation frequency was measured by parametric driving technique and found to be in excellent agreement with the predicted one. We keep the first order diffraction beam at its full power throughout the MOT loading time and atoms are loaded into the dipole trap mainly during the CMOT phase. Mechanical shutters are then used to block all resonant light from reaching the trapped atoms. The MOT pump beam, associated with the $|F = 2\rangle$ ground state, is blocked only a few milliseconds later to allow a 1 ms optical pumping pulse, effectively transferring all atoms into the $|F = 1\rangle$ ground state. Eventually, we are left with $\sim 10^6$ trapped atoms. Phase-space density conditions are similar to those resulted from the CMOT phase.

We note that the loading efficiency of the optical trap from the CMOT is rather poor due to the small spatial overlap between the two traps. Even for 100 W laser power the high initial temperature of the ^{7}Li atoms oblige tight focusing of the beam in order to create a deep enough potential. This makes the use of a compressible crossed dipole trap to improve the loading conditions impractical [5].

III. DETECTION OF FESHBACH RESONANCES ON $|F = 1, m_F = 0\rangle$ STATE

The scattering length of ^{7}Li atoms on all sublevels and sublevel mixtures of the $|F = 1\rangle$ ground state at zero magnetic field is positive but very small ($\sim 10a_0$), impeding efficient evaporation cooling. However, each sublevel and their mixtures posses at least one Feshbach resonance [18]. The resonance on the absolute ground state $|F = 1, m_F = 1\rangle$ has been previously used for the final stage of forced evaporation in an optical trap after precooling in a magnetic trap [19, 20]. The resonances on other sublevels or their mixtures were neither reported nor used before.

The search for Feshbach resonances requires high offset magnetic fields. We use two pairs of Helmholtz coils that allow variable bias fields of up to 1200 G. When the magnetic field is ramped up to high values we observe $\sim 50\%$ reduction in atom number. Our state selective measurement shows that the remaining atoms are all on $|F = 1, m_F = 0\rangle$ state. We attribute this loss to spin-flip collisions between atoms on $|F = 1, m_F = -1\rangle$ and $|F = 1, m_F = 1\rangle$ states. The energy level splitting diagram of the $|F = 1\rangle$ state is shown in Fig. 2. A spin-flip collision that takes place at high magnetic fields flips the colliding atoms to $|F = 1, m_F = 0\rangle$ state and leaves them with an excess of kinetic energy equal to 1.4 mK, which is comparable to the trap depth. Most of the atoms escape but we observe some increase in $|m_F = 0\rangle$ population. Detection of losses and spontaneous spin polarization are the first manifestations of the increase in collisional cross-section. As $|m_F = 0\rangle$ state is stable against spin-exchange collisions at high magnetic fields, we search for signatures of Feshbach resonances on this state.

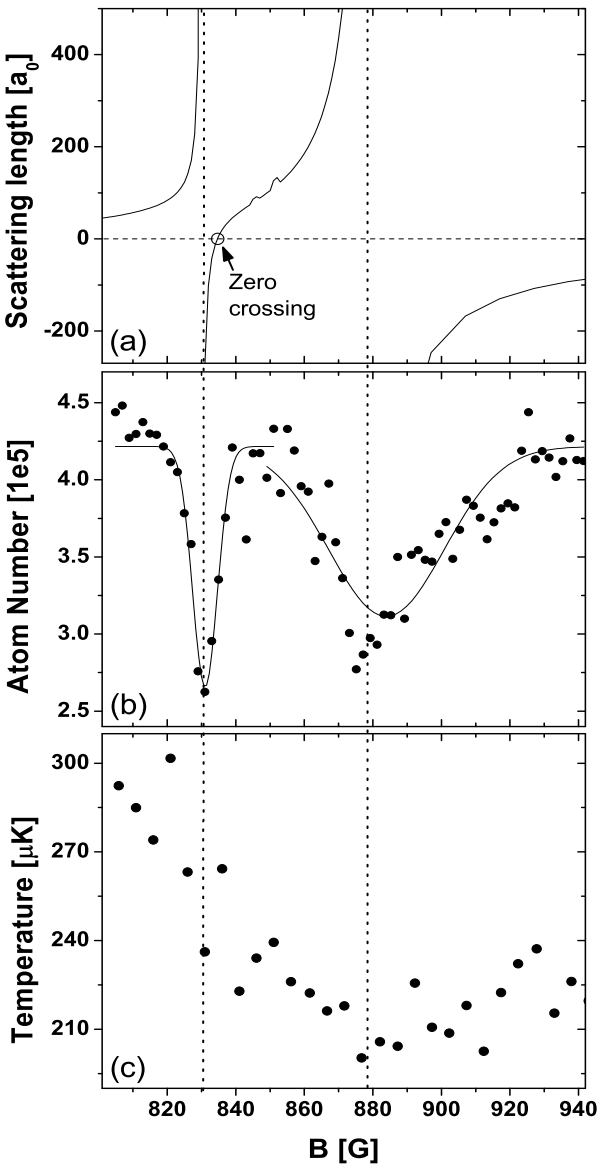

Figure 3: (a) Theoretical predictions for the scattering length as a function of bias magnetic field in the $|F = 1; m_F = 0\rangle$ state [18]. Divergence of the scattering length signify the presence of two Feshbach resonances. (b) The resonances are detected by measuring atom loss due to inelastic collisions. The solid lines are Gaussian fits to define resonances' positions. Note that the maximal loss does not coincide with the minimum of the gaussian fit for the broad resonance. (c) Temperature of the atomic cloud in the optical trap as a function of bias magnetic field. Decrease in temperature indicates cooling by free evaporation.

Fig. 3(a) shows a theoretical prediction of the scattering length as a function of bias magnetic field. Two Feshbach resonances between 800 G and 900 G are indicated by the divergence of the scattering length. Experimentally, the resonances are usually observed by detection of atom loss as in their proximity the inelastic collisional rate is strongly enhanced [21, 22]. For this measurement we ramp up the magnetic field to different values in 40 ms and wait for 0.5 s before decreasing it to 386 G in 40 ms where *in situ* state selective absorption imaging is performed. In Fig. 3(b) the atom number as a function of bias magnetic field is shown. The atoms are initially prepared at a temperature of 300 μK. Strong losses are observed around 830 G and 880 G featuring a narrow and a broad structures. By fitting them with simple Gaussian functions, a coarse estimation of the resonances locations and widths can be determined. According to the fit, the narrow resonance is 7 G wide ($1/e^2$ radius) and it is located at 831 ± 4 G. The uncertainty in its position is due to an uncertainty in the magnetic field calibration which is obtained by the detection of optical resonances and thus limited by the linewidth of the excited states and the laser locking quality. The broad resonance located at 884^{+4}_{-13} G is 34 G wide and features a notable asymmetric profile which tends to shift the center of a simple gaussian fit to a higher magnetic field value as compared to the maximal loss position (detected at 875 G). Such an asymmetry in losses in the vicinity of broad Feshbach resonances has been recently reported for ^{39}K atoms [23] and was attributed to larger three-body loss coefficient from the negative scattering length side of the resonance and mean field effects. The followed study of molecule association showed that the position of a broad Feshbach resonance is indeed shifted to a lower value [23]. Comparison between the experimental measurements (Fig. 3(b)) and the theoretical calculations (Fig. 3(a)) shows that the maximal loss position suits better the predicted distance between the two resonances. We therefore believe that the actual location of the broad resonance is at somewhat lower value than that given by the Gaussian fit and this systematic error increases the error bars of the experimentally detected resonance position.

Most excitingly, we observe an onset of free evaporation cooling together with the enhanced inelastic losses (Fig. 3(b)). In Fig. 3(c) we show the temperature change as a function of the bias magnetic field. A decrease in the temperature indicates cooling which we attribute to the establishment of free evaporation when the collision rate becomes high enough. The observed cooling allows us to decrease temperature by forced evaporation and than scan for Feshbach resonances again to improve sensitivity.

We execute a short forced evaporation of 1.5 s reducing the trap depth to 0.3 mK, 15% of its initial value. Evaporation is performed with a bias field of 850 G which is chosen to optimize number-to-temperature ratio based on our scans for Feshbach resonances at high temperature (Fig. 3(b,c)). By the end of this evaporation the atoms are cooled down to a temperature of $\sim$50 μK. Our scan for inelastic losses in proximity to Feshabch resonances reveals the same positions of the resonances observed before including asymmetry in the profile of the broad resonance. However, an additional feature has been identified which we were unable to detect at high temperature.

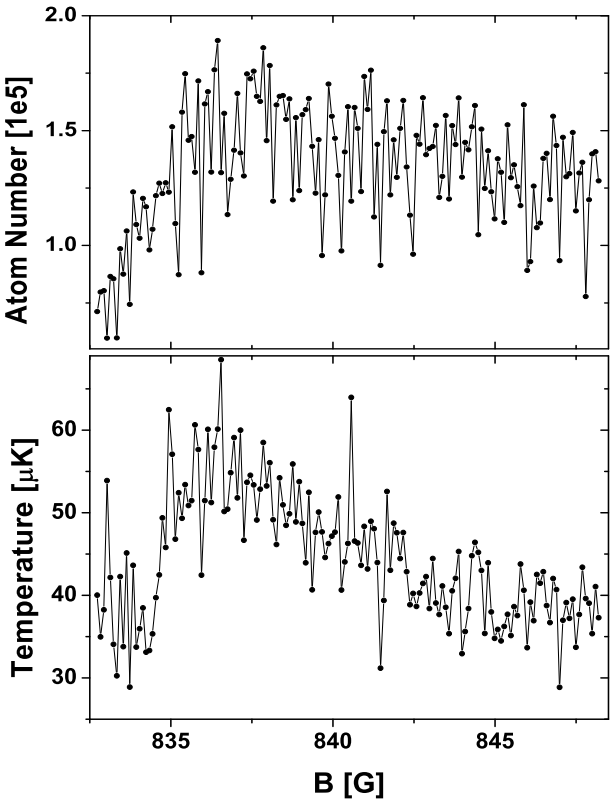

Figure 4: The gain in atom number and increase in temperature, which indicate the lack of free evaporation cooling, reveal the zero crossing of the scattering length at 836 ± 4 G.

In Fig. 4, zero crossing of the scattering length manifests itself in terms of increased temperature and number of atoms [24]. The hold-time at high bias field for this experiment is 0.5 s. During this time efficient free evaporation reduces the temperature to 40 μK if collisional cross-section is high enough. At zero crossing the scattering length vanishes and so does the cross-section, impeding efficient free evaporation. The asymmetry in this measurement is caused by the proximity of the zero crossing point to the narrow resonance at 831 G. The zero-crossing point is detected at 836 ± 4 G where the maximum temperature is observed.

IV. EVAPORATION COOLING TO THE BEC THRESHOLD

Forced evaporation cooling of the optically trapped atoms down to the BEC threshold is performed by attenuating the laser power, thus reducing the trap depth which scales linearly with the power reduction factor α. A photo diode detector, located behind one of the mirrors, collects a fraction of the trap beam's power (see Fig. 1). It generates a voltage readout that is compared to a setpoint signal by a PID-controller. An error signal is fed back to the RF-power supplier of the AOM. With

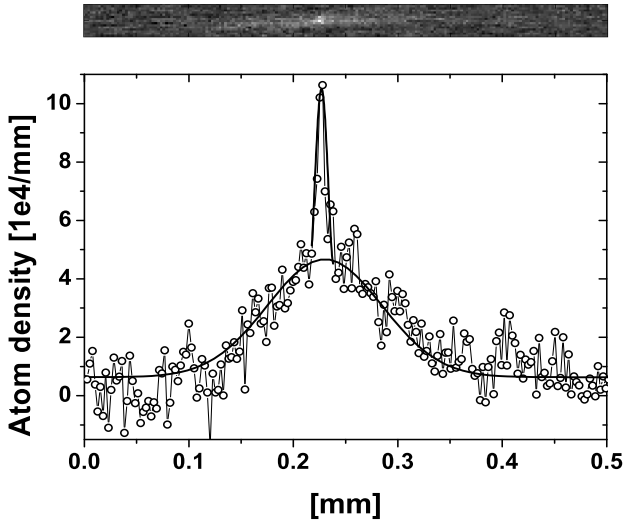

Figure 5: On-set of BEC. *In-situ* absorption imaging of atoms at the BEC threshold. The thermal atomic cloud is fitted with a Bose-Einstein distribution function which yields the temperature of 380 ± 40 nK. The number of atoms in the BEC is ~ 700.

this scheme we are able to control trap depth reduction up to a factor of $\alpha = 3 \times 10^{-3}$. As is well known, trap oscillation frequencies also decrease with the attenuation of laser power ($\propto \sqrt{\alpha}$), affecting re-thermalization efficiency throughout the evaporation process. In addition to that, the strong bias magnetic field, employed during evaporation, creates a weak anti-trapping potential in the longitudinal direction. This decreases further the optical confinement toward the end of the evaporation. Indeed, in a single beam trap we were unable to cool atoms below a temperature of 10 μK. The addition of the zero-order diffraction beam solves this problem. At the beginning of the evaporation it has no effect on the atoms as its potential depth is negligible. However, with the reduction of the first order diffraction intensity, it strengthen to create a confinement potential to the atoms in the longitudinal direction of the trap. The longitudinal oscillation frequency, determined by the confining beam, is $2\pi \times 60$ Hz. The zero order beam can be effectively considered as a two dimensional confining potential because it produces very weak (0.14 Hz) oscillation frequency in its propagation direction which is easily overcome by the magnetic anti-trapping potential. We note also that the crossed trap is slightly shifted from the location of the zero order beam waist which reduces somewhat the potential at the end of the evaporation.

Forced evaporation is found to work most effectively when the bias field is set to 866 G. Based on the theoretical curve (Fig. 3(a)) we estimate the scattering length to be $(300 \pm 100)a_0$. The large uncertainty in the scattering length is due to the uncertainty in the position of the broad resonance. The trap depth is reduced exponentially in 3 s with a time constant of 330 ms to less

than 0.5% of its initial value. In Fig. 5, *in-situ* absorption imaging of the trapped atoms at 866 G reveals the on-set of a BEC threshold by a familiar bi-modal density distribution. The trace represents the atom longitudinal density after integrating the radial direction of the picture above it. Optical resolution is 4 μm, less than the size of the BEC. The thermal atomic cloud is fitted with a Bose-Einstein distribution function which reveals a temperature of $T = 380 \pm 40$ nK. The total number of atoms is 6×10^3 which sets the critical temperature to $T_c = 350$ nK. The fitting of the BEC with the inverted parabola of a Thomas-Fermi limit reveals ~ 700 atoms in the condensate.

V. CONCLUSIONS

To conclude, we developed a method to all-optically condense ^{7}Li atoms. This way facilitates the BEC production which is extremely demanding in magnetic traps. We observed a BEC with repulsive interactions on $|F = 1, m_F = 0\rangle$ state in less than 3 s of forced evaporation in a crossed beam optical dipole trap. We use the tunability of the interatomic interactions in the proximity of Feshbach resonances which we observed and characterized for this internal state.

We note that the condensate life-time was very short, presumably because of the extremely large scattering length. Moreover, our weak optical trap at the end of the evaporation was not stable enough due to the use of a single linear photo diode detector. A number of improvements can be implemented to optimize the performance of the described method. Better laser beam stability, which can be achieved by using either two detectors [7] or a logarithmic detector, and decrease of the scattering length toward the end of evaporation will improve the condensate lifetime. Improved vacuum would allow an increase in the number of atoms in the condensate.

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